AMENDMENTS TO THE CLAIMS

The listing of claims will replace all prior versions, and listings, of claims in the specification:

Listing of the Claims:

1. (Currently Amended) A carrier with a non-cationic surface, which can accumulate on a damaged endothelial cell site of a tissue comprising endothelial cells, wherein the carrier comprises a carboxylic type lipid that has no phosphate group.

2. (Original) A carrier according to claim 1, wherein the surface is a membrane.

3. (Original) A carrier according to either one of claims 1 and 2, wherein the tissue is a vessel.

4. (Original) A carrier according to claim 3, which can diffuse outside the vessel.

5. (Previously Presented) A carrier according to claim 3, wherein the vessel is a blood vessel.

6. (Original) A carrier according to claim 1, wherein the damage reaches an endothelial cell.

7. (Original) A carrier according to claim 1, wherein the damage comprises those that result from laser, inflammation, ischemic disorder, ischemia-reperfusion damage, bacterial toxin, oxidative stress, tumor or thrombus formation, or bleeding.

8. (Original) A carrier according to claim 7, wherein the inflammation is brain edema.

9. (Original) A carrier according to claim 7, wherein the ischemic disorder is cerebral ischemic disorder.

10. (Original) A carrier according to claim 7, wherein the ischemia-reperfusion damage is ischemia-reperfusion-induced organ damage.

11. (Cancelled)

12. (Currently Amended) A pharmaceutical composition comprising the ~~drug transporter of~~ carrier according to claim ~~11~~ 1 incorporating or carrying a drug.

13. (Original) A pharmaceutical composition according to claim 12, which functions as a drug for controlling a platelet function.

14. (Original) A pharmaceutical composition according to claim 13, wherein the platelet function to be controlled comprises hemostasis, antithrombotic formation, thrombolysis or antiatherogenic action.

15. (Original) A pharmaceutical composition according to claim 12, wherein the drug is at least one selected from a group consisting of substances that are activated by light, change in temperature, change in pH, ultrasound, uptake of an inflammation-mediating cell or enzyme degradation; hemostatic agents; antithrombotic agents; thrombolytic agents; antitumor agents; and antiatherogenic agents.

16. (Original) A pharmaceutical composition according to claim 15, wherein the inflammation-mediating cell is a lymphocyte, a leukocyte, a macrophage or a platelet.

17. (Currently Amended) A drug delivery method comprising *in vivo* administering the pharmaceutical composition of claim 12 and allowing ~~the pharmaceutical~~ said composition ~~according to claim 12~~ to accumulate on a damaged site of a tissue.

18. (Previously Presented) A drug control method comprising allowing the pharmaceutical composition according to claim 12 to accumulate on a damaged site of a tissue and allowing the drug to act on the damaged site.

19. (Original) A method according to claim 18, wherein the action of the drug is controlled by accumulation of the carrier, diffusion of the carrier or activation of the carrier.

20. (Original) A method according to any one of claims 17 to 19, wherein the tissue is a vessel.

21. (Original) A method according to claim 20, wherein the vessel is a blood vessel.

22.-27. (Cancelled)